PATENT COOPERATION TREATY



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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

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INTERNAT	TIONAL PRELIMINAR	Y EXAMIN	ATION REP	ORT
	(PCT Article 36 a	nd Rule 70)		
Applicant's or agent's file reference PJ644-113PCT	FOR FURTHER ACTIO	N See Notifi Preliminary	cation of Tra Examination Re	nsmittal of International port (Form PCT/IPEA/416)
international application No. PCT/FR2003/002091	International filing date (da 04 juillet 2003 (04			day/month/year) bre 2002 (11.12.2002)
nternational Patent Classification (IPC) o C12Q 1/68	r national classification and IPC			
Applicant CENTRE 1	NATIONAL DE LA REC	HERCHE SC	IENTIFIQUE	3
amended and are the basis 70.16 and Section 607 of the	t according to Article 36.	iding this cover a s of the description attaining rectification of the PCT).	sheet.	or drawings which have beer
This report contains indications r	relating to the following items:			
I Basis of the repo	ort			
II Priority				
III Non-establishme	ent of opinion with regard to not	elty, inventive s	tep and industria	al applicability
IV Lack of unity of		-14. :	-Marie -Marie an	' '
V Reasoned statem citations and exp	nent under Article 35(2) with repolarized such states	ard to novelty, inent	nventive step or	industriai applicaoility,
VI Certain documer	nts cited			
VII Certain defects i	n the international application			
VIII Certain observat	tions on the international applica	tion		
Date of submission of the demand	Da	te of completion	of this report	
15 juin 2004 (15.0	6.2004)	22 I	February 2005	5 (22.02.2005)
Name and mailing address of the IPEA/	EP Au	thorized officer		
Racsimile No	To	enhone No.		

International application No.

PCT/FR2003/002091

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

L. B	asis o	of the re	port	
1. \	With 1	regard to	the elements of the international application:*	
	X	the inte	mational application as originally filed	
Ī	$\overline{\boxtimes}$	the desc	ription:	1
		pages	1-24	, as originally filed
		pages		, filed with the demand
		pages	, filed with the letter of	
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1	\triangle			, as originally filed
		pages	, as amended (togethe	
ļ		pages	,	, filed with the demand
		pages	1-19 , filed with the letter of	10 September 2004 (10.09.2004)
וו	Δ	the drav		, as originally filed
1		pages	1/20-20/20	, filed with the demand
		pages	, filed with the letter of	
١,		pages		
	t	he seque	nce listing part of the description:	1
		pages		, as originally filed
Į		pages		, filed with the demand
		pages	, filed with the letter of	
	the in	nternation e elemen	, , , , , , , , , , , , , , , , , , , ,	which is:
		the lan	guage of a translation furnished for the purposes of international search (under I	Rule 23.1(b)).
	Щ		guage of publication of the international application (under Rule 48.3(b)).	
	Ш	the lar or 55.3	guage of the translation furnished for the purposes of international preliminals).	ry examination (under Rule 55.2 and/
3.	With	n regard minary e	to any nucleotide and/or amino acid sequence disclosed in the intern examination was carried out on the basis of the sequence listing:	ational application, the international
		contai	ned in the international application in written form.	
ı	Ш	filed to	ogether with the international application in computer readable form.	
ļ		furnisl	ned subsequently to this Authority in written form.	
			ned subsequently to this Authority in computer readable form.	
			tatement that the subsequently furnished written sequence listing does national application as filed has been furnished.	ot go beyond the disclosure in the
	Ш		tatement that the information recorded in computer readable form is identic turnished.	al to the written sequence listing has
4.		The ar	mendments have resulted in the cancellation of:	
			the description, pages	
1		\Box	the claims, Nos.	
1			the drawings, sheets/fig	
5.		This re	eport has been established as if (some of) the amendments had not been made, I the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**	since they have been considered to go
	in th	acement his repoi 70.17).	sheets which have been furnished to the receiving Office in response to an inv rt as "originally filed" and are not annexed to this report since they do	itation under Article 14 are referred to not contain amendments (Rule 70.16
**	Any	replacen	nent sheet containing such amendments must be referred to under item 1 and an	nexed to this report.

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V.	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability;
	citations and explanations supporting such statement

1. Statement			
Novelty (N)	Claims	4-19	YES
	Claims	1-3	NO NO
Inventive step (IS)	Claims		YES
	Claims	1-19	NO
Industrial applicability (IA	L) Claims	1-19	YES
	Claims		NO NO

2. Citations and explanations

- 1.1 Reference is made to the following documents:
 - D1: US-A-4 238 757 (SCHENK JOHN F) 9 December 1980 (1980-12-09)
 - D2: SOUTEYRAND E ET AL: "DIRECT DETECTION OF THE HYBRIDISATION OF SYNTHETIC HOMO-OLIGOMER DNA SEQUENCES BY FIELD EFFECT" JOURNAL OF PHYSICAL CHEMISTRY. B, MATERIALS, SURFACES, INTERFACES AND BIOPHYSICAL, WASHINGTON, DC, US, vol. 101, 1997, pages 2980-2985, XP001040796 ISSN: 1089-5647
 - D3: TSURUTA H. ET AL: "Detection of the products of a polymerase chain reaction by an ELISa system based on an ion sensitive field effect transistor" JOURNAL OF IMMUNOLOGICAL METHODS, vol. 176, 1994, pages 45-52, XP009021947
 - D4: WO 03/054225 A (BIOCHIP TECHNOLOGIES GMBH; LEHMANN MIRKO (DE); MICRONAS GMBH (DE);) 3 July 2003 (2003-07-03)
 - D5: WO 03/052097 A (MIYAHARA YUJI; HATTORI KUMIKO (JP); YASUDA KENJI (JP); HITACHI HIG) 26 June 2003 (2003-06-26)
- 1.2 Documents D4 and D5 have an earlier priority date

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and/or filing date than the present international patent application. Therefore, the subject matter of these documents may be relevant as far as the present international patent application is concerned in the regional or national phase.

2. Novelty

Document D1 describes a field effect transistor 2.1 wherein the active transistor region ("gate") is loaded with antibodies. When said transistor is contacted with a buffer containing the antigen specific to said antibody, interaction between the antibody and the antigen can be measured using a source/drain current. Measurement can be enhanced by reducing the salt concentration in the measurement buffer relative to the salt concentration in the interaction buffer between the antibody and the antigen (see D1, the abstract; figures 1 and 2; column 3, line 15 to column 6, line 45). D1 also discloses how a plurality of substances can be measured simultaneously (see D1, figure 3; column 5, line 42 to column 6, line 23) and how the measurements are carried out differentially (see D1, column 4, lines 11-39).

The subject matter of the claims is defined in terms of the differential measurement method.

However, no additional specific information on measurements is indicated in the claims (e.g. a measurement carried out in the time-dependent state or the steady state). It follows that the subject matter of claims 1 to 3 lacks novelty (PCT Article 33(2)).

2.2 The amendments submitted with the letter of 10

September 2004 do not cause the subject matter of the application to be extended beyond the content of the application as filed. Therefore, they are consistent with the provisions of PCT Article 34(2)(b).

3. Inventive step

3.1 Dependent claims 4 to 19 do not contain any features which, when combined with the features of any one of the claims to which they refer, might define subject matter that complies with the requirements of inventive step of the PCT, in the light of document D2, which describes the direct detection of oligonucleotide hybridisation using a field-effect transistor (D2, title; abstract; page 2981, figure 1; page 2983, figure 3, page 2985, figure 11) and document D3, which describes the detection of the products of a polymerase chain reaction using a field-effect transistor (D3, title; abstract; page 47, figure 1; page 50, figures 3 and 4).

4. Industrial applicability

4.1 The subject matter of the application, as defined in claims 1 to 19, appears to be industrially applicable (PCT Article 33(4)).